Product Datasheet

Mbo II

5'...**GAAGA(N)**₈...3' 3'...CTTCT(N)₇...5'

Product No : RE1290 Quantity : 100u



Lot **Expiry Date**

Concentration 5u/ul 1ml of 10X Buffer Mbo II Supplied with

1ml of 10X Buffer UB 0.5ml Diluent Viva Buffer A

(BSA included in all Reaction Buffer)

Store at -20°C



info@vivantechnologies.com

Reaction Conditions:

Buffer Mbo II,

30mM Tris-acetate (pH 7.9 at 30°C), 10mM Mg-acetate, 60mM K-acetate, 7mM 2-mercaptoethanol and 100µg/ml BSA. Incubate at 37°C.

Dilution: Viva Buffer A

10mM Tris-HCI (pH 7.4 at 25°C), 50mM KCI, 0.1mM EDTA, 1mM DTT, 200µg/ml BSA and 50% glycerol.

Thermal Inactivation: 65°C for 20 minutes

Storage Buffer:

10mM Tris-HCI (pH 7.5), 100mM NaCl, 0.1mM EDTA, 7mM 2-mercaptoethanol, 100µg/ml BSA and 50% glycerol.

Unit Definition:

1u is defined as the amount of enzyme that is required to digest 1μg of DNA in 1 hour at 37°C in 50μl of assay buffer.

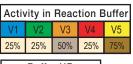
Quality Control Assays:

Ligation/ Recutting Assay:

After 5-fold overdigestion with Mbo II, more than 90% of the DNA fragments can be ligated and recut.

Overdigestion assay:

An unaltered banding pattern was observed after 1μg of DNA was digested with 10u of Mbo II for 16 hours at 37°C (Without BSA).





* Buffer UB is provided for double digestion purpose.

NOTE:

- *Blocked by overlapping dam-methylation (GmATC): **GAAGA**TC.
- * Total reaction volume dependent on experiment.
- * The amount of enzyme to be used is very much dependent on the DNA template.
- * For plasmid DNA, 5-10X more enzyme is required.

Example of Digestion Reaction

: 1 unit Enzvme

Lambda(dam- & dcm-)0.3µg/µl : 3.33µl (1µg DNA)

10X Reaction Buffer : 5µl

Sterile Distilled Water : Up to 50µl

Product Use Limitation

This product is for research purposes and in vitro use only. V I V a n t I S | www.vivantechnologies.com